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(54) Title: USE OF 13C–NMR TO DETECT BINDING

(57) Abstract

Methods of detecting binding of a putative ligand to a ^{13}C–enriched target molecule, methods of screening for compounds which bind to a ^{13}C–enriched target molecule, methods for calculating the dissociation constant of a ligand compound which binds to a ^{13}C–enriched target molecule, and methods employed in the determination of the specific amino acids in a ^{13}C–enriched target molecule affected by the binding of a ligand, as well as compounds identified by these screening methods, are provided herewith.

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USE OF ^{13}C–NMR TO DETECT BINDING

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Field of the Invention

The present invention relates to the use of nuclear
magnetic resonance to detect binding between compounds and ^{13}C-
10 enriched target molecules.

Cross-Reference to Related Applications

15   This application is a continuation-in-part of application
Serial No. 09,241,194, filed February 1, 1999, which is a
continuation of application Serial No. 08/744,701, filed
October 31, 1996, now United States Patent No. 5,---,---,
which is a continuation-in part of application Serial No.
20 08/678,903, filed July 12, 1996, now abandoned, which is a
continuation-in-part of application Serial No. 08/558,633,
filed November 14, 1995, now United States Patent 5,---,---.

25   ### Background of the Invention

One of the most powerful tools for discovering new drug
leads is random screening of synthetic and natural product
libraries to discover compounds that bind to a particular
30 target molecule (for example, the identification of ligands to

that target). Using this method, ligands may be identified by
their ability to form a physical association with a target
molecule or by their ability to alter a function of a target
molecule.

5   When physical binding is sought, a target molecule is
typically exposed to one or more compounds suspected of being
ligands and assays are performed to determine if complexes
between the target molecule and one or more of those compounds
are formed. Such assays, as is well known in the art, test

10   for gross changes in the target molecule (for example, changes
in size, charge, mobility) that indicate complex formation.

      Where functional changes are measured, assay conditions
are established that allow for measurement of a biological or
chemical event related to the target molecule (for example, an

15   enzyme-catalyzed reaction, receptor-mediated enzyme
activation, and the like). To identify an alteration, the
function of the target molecule is determined before and after
exposure to the test compounds.

      Existing physical and functional assays have been used

20   successfully to identify new drug leads for use in designing
therapeutic compounds. There are, however, limitations
inherent to those assays that compromise their accuracy,
reliability and efficiency. A major problem with existing
assays, for example, relates to the generation of "false

25   positives". In a typical functional assay, a "false positive"
result is generated for a compound that triggers the assay but
which compound is not effective in eliciting the desired
physiological response. In a typical physical assay, a "false
positive" is a compound that, for example, attaches itself to

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the target but in a nonspecific manner (for example,
non-specific binding). False positives are particularly
prevalent and problematic when screening higher concentrations
of putative ligands because many compounds have non-specific
5   effects at those concentrations.

In a similar fashion, existing assays are frequently
plagued by the problem of "false negatives", which result when
a compound gives a negative response in the assay but, as
found subsequently by some other method, is actually a ligand
10   for the target. False negative results typically occur in
assays that use concentrations of test compounds that are
either too high (resulting in toxicity) or too low, relative
to the binding or dissociation constant of the compound to the
target.

15   Another problem with existing assays is the limited
amount of information that is provided by the assay itself.
While the assay may correctly identify compounds that attach
to or elicit a response from the target molecule, those assays
typically do not provide any information about either specific
20   binding sites on the target molecule or structure activity
relationships between the compound being tested and the target
molecule. The inability to provide any such information is
particularly problematic where the screening assay is being
used to identify leads for further study.

25   It has recently been suggested that X-ray crystallography
can be used to identify the binding sites of organic solvents
on macromolecules. However, this method cannot determine the
relative binding affinities at different sites on the target.
It is only applicable to very stable target proteins that do

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not denature in the presence of high concentrations of organic solvents. In addition, due to the long time needed to determine the individual crystal structures, this approach is not a suitable method for rapidly testing a large number of

5   compounds that are chemically diverse, but is limited to mapping the binding sites of only a few organic solvents.

Rapid, efficient, and reliable methods of determining ligand/target binding, and mapping binding sites on the target substance are disclosed in United States Patent Nos. 5,698,401

10   and 5,804,390, to Fesik, et al. These patents disclose methods of detecting binding of a ligand compound to a target biomolecule by generating first and second nuclear magnetic resonance correlation spectra from target biomolecules which have been isotopically enriched with the NMR-detectable ^{15}N

15   nucleus. The first spectrum is generated from data collected on the target substance in the absence of ligands, and the second in the presence of one or more ligands. A comparison of the two spectra permits determination of which compounds in the mixture of putative ligands bind(s) to the target

20   biomolecule, as well as specific information about the site of binding. Since the methods elicit information about the binding sites on the target molecule, the methods can be used for optimizing the design of ligands to a pre-selected target.

Because the NMR methods of Fesik, et al., supra, require

25   nitrogen-containing target substances due to the dependence of those methods on isotopic-enrichment of the target molecule with ^{15}N, it would be a valuable contribution to the art to provide an alternative method which employs a different type of isotopically-enriched target molecule.

5

Summary of the Invention

The instant invention provides a method of detecting

5   binding between one or more putative ligands to a pre-
selected, isotopically-enriched target molecule.

The instant invention further provides a method of
screening a mixture of compounds for binding to a pre-
selected, isotopically-enriched target molecule.

10   Also provided by the instant invention is a method of
determining the dissociation constant for a ligand compound
that binds to a pre-selected, isotopically-enriched target
biomolecule.

Additionally provided by the instant invention are

15   methods employed in the determination of the specific amino
acids in a ^{13}C-enriched target molecule affected by the binding
of a ligand.

Still further provided by the instant invention is a
compound identified by a method of screening a mixture of

20   compounds for binding to a pre-selected, isotopically-enriched
target molecule.

Brief Description of the Drawings

25

Figure 1 illustrates a ^{13}C/^{1}H correlation spectrum of
uniformly ^{13}C-labeled FKBP.

Figure 2 illustrates the methyl regions of ^{13}C/^{1}H
correlation spectra of uniformly ^{13}C-labeled FKBP before (thin

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multiple contours) and after (thick single contours) addition of 2-   phenylimidazole (0.12 mM).

Figure 3 illustrates the methyl regions of $^{13}C/^1H$ correlation spectra of FKBP selectively $^{13}C/^{15}N/^2H$-labeled at valinyl and leucyl residues before (thin multiple contours) and after (thick single contours) addition of 2-phenylimidazole (0.25 mM).

Figure 4 illustrates the methyl regions of $^{13}C/^1H$ correlation spectra of FKBP selectively ^{13}C-labeled at alanyl residues before (thin multiple contours) and after (thick single contours) addition of 2-phenylimidazole (0.25 mM).

Figure 5 illustrates a "stick model" depiction of the three-dimensional structure of FKBP.

Detailed Description of the Invention

Terms used throughout this specification have their usually accepted meanings. The following specific terms have the ascribed meanings.

"DTT" means dithiothreitol.

"FKBP" refers to FK-binding protein.

"HEPES" denotes N-2-hydroxyethylpiperazine-N'-2-ethylsulfonic acid.

"IPTG" means isopropyl-β-D-thiogalactopyranoside.

"PMSF" refers to α-toluenesulfonyl fluoride.

"SCD" refers to the catalytic domain (residues 81-256) of stromelysin.

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Any target biomolecule which gives a high resolution NMR spectrum and which can be uniformly or selectively enriched with ^{13}C can be used in the methods of the present invention. These methods are thus applicable to any desired ^{13}C-enriched

5   target biomolecule, including lipoproteins, lipoprotein fragments, glycoproteins, glycoprotein fragments, proteins, protein fragments, polypeptides, DNA and RNA. The natural isotopic abundance of ^{13}C is 1.11%. Thus, the probability that any given carbon atom within an organic molecule is ^{13}C is

10 normally about 0.0111.

In order to increase the strength of a NMR signal evincing data related to spin coupling between the nucleus of a ^{13}C carbon atom and any adjacent hydrogen atoms, it is desirable to increase the natural ^{13}C content of the target

15 molecule being studied. This is accomplished by either uniformly or selectively enriching the target molecule with ^{13}C. As used throughout this specification and the appended claims, the terms "uniform enrichment," "uniformly enriching," "uniformly enriched," uniform labeling" and "uniformly

20 labeled" mean increasing to a value greater than 0.0111, by synthetic means, the probability that a carbon atom randomly selected throughout the target molecule will be ^{13}C. On the other hand, the terms "specific enrichment," "specifically enriching," "specifically enriched," "specifically labeling"

25 and "specifically labeled" mean increasing to a value greater than 0.0111, by synthetic means, the probability that carbon atoms at one or more specific pre-selected site(s) within the target molecule will be ^{13}C.

8

For example, biomolecules expressed by genetically modified microorganisms grown in a nutrient medium containing uniformly ^{13}C-enriched glucose will be uniformly ^{13}C-enriched. On the other hand, a protein expressed by a genetically

5   modified microorganism in a nutrient medium containing alanine that is ^{13}C-enriched only on the methyl side chain will be specifically enriched by ^{13}C at those alanyl residues contained within the expressed protein.

The method of the present invention, which employs

10   isotopic enrichment with ^{13}C, is applicable to any organic target molecule, including those containing nitrogen. The method also permits the analysis of target molecules in which specific carbon atom sites have been enriched (for example, methyl groups of alanyl, leucyl, isoleucyl, and valinyl

15   residues). Methyl groups have favorable relaxation properties compared to amide groups, which is advantageous when applied to larger target biomolecules (MW > 30 kDa).

Polypeptides and proteins perform many pivotal roles in living organisms. The examples provided below employ

20   polypeptides to illustrate the instant method. Polypeptides and protein fragments comprise preferred classes of target substances for the method of the present invention. However, it is to be understood that the method of the present invention is applicable to other target substances which can

25   be ^{13}C-enriched.

The preparation of uniformly and specifically ^{13}C-enriched exemplary polypeptide target molecules is set forth below. One means of preparing adequate quantities of either uniformly or specifically ^{13}C-enriched polypeptide-containing target

molecules involves the transformation of a host cell with an
expression vector containing a polynucleotide encoding the
desired polypeptide. The protein or polypeptide protein
fragment is expressed by culturing the transformed cell line
5   in a medium containing assimilable sources of ^{13}C well known in
the art. A preferred assimilable source for uniform ^{13}C
labeling is uniformly ^{13}C-labeled glucose or U-^{13}C-glucose,
available from Cambridge Isotope Laboratories. For site-
specific labeling, assimilable sources for ^{13}C-labeling of a
10 target polypeptide include commercially available specifically
^{13}C-labeled amino acids. In an alternative embodiment for
specific enrichment, the assimilable sources of ^{13}C contained
in the nutrient medium are ^{13}C-labeled biosynthetic precursors
of amino acids. For example, α-keto-butyrate is the
15 biosynthetic precursor of isoleucine and α-keto-isovalerate is
the biosynthetic precursor of both valine and leucine. Scheme
I below shows how the specifically ^{13}C-enriched biosynthetic
precursors of leucine, isoleucine, and valine, can be
synthesized. The comparatively inexpensive ^{13}C-enriched methyl
20 iodide ($H_3^{13}CI$) may be employed as the source for isotopic
enrichment to produce C-terminally-labeled α-keto-butyric acid
and α-keto-isovaleric acid.

The use of a uniformly ^{13}C-enriched nutrient such as ^{13}C-
enriched glucose is a convenient means of introducing ^{13}C-
25 enrichment into a target substance; however, it is very
expensive. Furthermore, a vast majority of the carbon sites
in uniformly ^{13}C-labeled targets will have a covalently bonded
neighbor which is also ^{13}C-labeled, introducing ^{13}C-^{13}C coupling
which can negatively impact both the signal-to-noise and

relaxation properties of ^{13}C-labeled sites in the target
biomolecule. A particular advantage is achieved by site-
specifically labeling the target polypeptide with [13]C. As
stated above, this can be accomplished by including

5   commercially available ^{13}C-labeled amino acids in the nutrient
medium. This too, however, is a costly alternative, but may
be desirable in some circumstances when labeling of certain
types of aminoacyl residues in the target polypeptide is
required. However, the preferred method of ^{13}C-labeling a

10 polypeptide target molecule comprises growing the genetically
modified cell line in a nutrient medium containing ^{13}C-labeled
biosynthetic precursors of amino acids. In particular,
preferred amino acid precursors that are labeled include α-
keto-butyric acid and α-keto-isovaleric acid. The

15 biosynthetic products of these precursors are leucine,
isoleucine, and valine in which particular side-chain methyl
groups are ^{13}C-enriched. Because the methyl groups each have
three hydrogen atoms connected to a ^{13}C-labeled carbon atom,
the corresponding NMR signals are particularly strong and

20 distinctive.

The synthetic sequence for labeled α-keto-butyric acid
and α-keto-isovaleric acid involves the methylation of the
terminal carbon atom in pyruvic acid with ^{13}C-enriched methyl
iodide. Normally, the alkylation of α-keto acids, such as

25 pyruvate, is inherently difficult and is accompanied by
decomposition of the enolate intermediate with the formation
of numerous side products. However, T. Spencer, et al.,
Tetrahedron_Letters, 1975, 3889, and D. R. Williams, et al.,

ibid, 1990, 5881, have shown that alkylation of the corresponding oxime can be easily accomplished, and D. Enders, *et al., Angew. Chem. Int. Eng. Ed.*, 1992, 618 and D. Enders, *et al., Synlett*, 1992, 901 have demonstrated that the

5   hydrazone is also readily alkylated without decomposition.

In Scheme I, *tert*-butyl pyruvate, **1**, is converted to the corresponding N,N-dimethylhydrazone, **2**, by reaction with N,N-dimethylhydrazine in diethyl ether at room temperature. The resulting hydrazone, **2**, is cooled in tetrahydrofuran solution

10   to -78°C, and treated with lithium bromide, followed by lithium diisopropylamide to form the intermediate aza-allyl enolate. The enolate is alkylated with ^{13}C-labeled methyl iodide to produce hydrazone **3**. A second course of alkylation of **3** produces the labeled dimethylated hydrazone, **4**. Treatment of

15   **3** and **4** with trifluoroacetic acid in methylene chloride at 0°C cleaves both the hydrazine and ester groups to produce the corresponding ^{13}C-terminally labeled α-ketoacids, **5** and **6** (4-(^{13}C)-butyric acid and 4-(^{13}C)-3-(^{13}C)-methylbutyric acid, respectively).

20

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Scheme I

Chemical Synthesis of Labeled Precursors

5

1 → N,N-Dimethylhydrazine / Cyclohexane, heat → **2**

1. (optional) LiBr, LDA
2. H_3^nCl, THF, -78°C

3

1. 1N aq HCl/THF or Et_2O
2. HCl gas/CH_2Cl_2

→ **5**

1. (optional) LiBr, LDA
2. H_3^*Cl, THF, -78°C

4

1. 1N aq HCl/THF or Et_2O
2. HCl gas/CH_2Cl_2

→ **6**

13
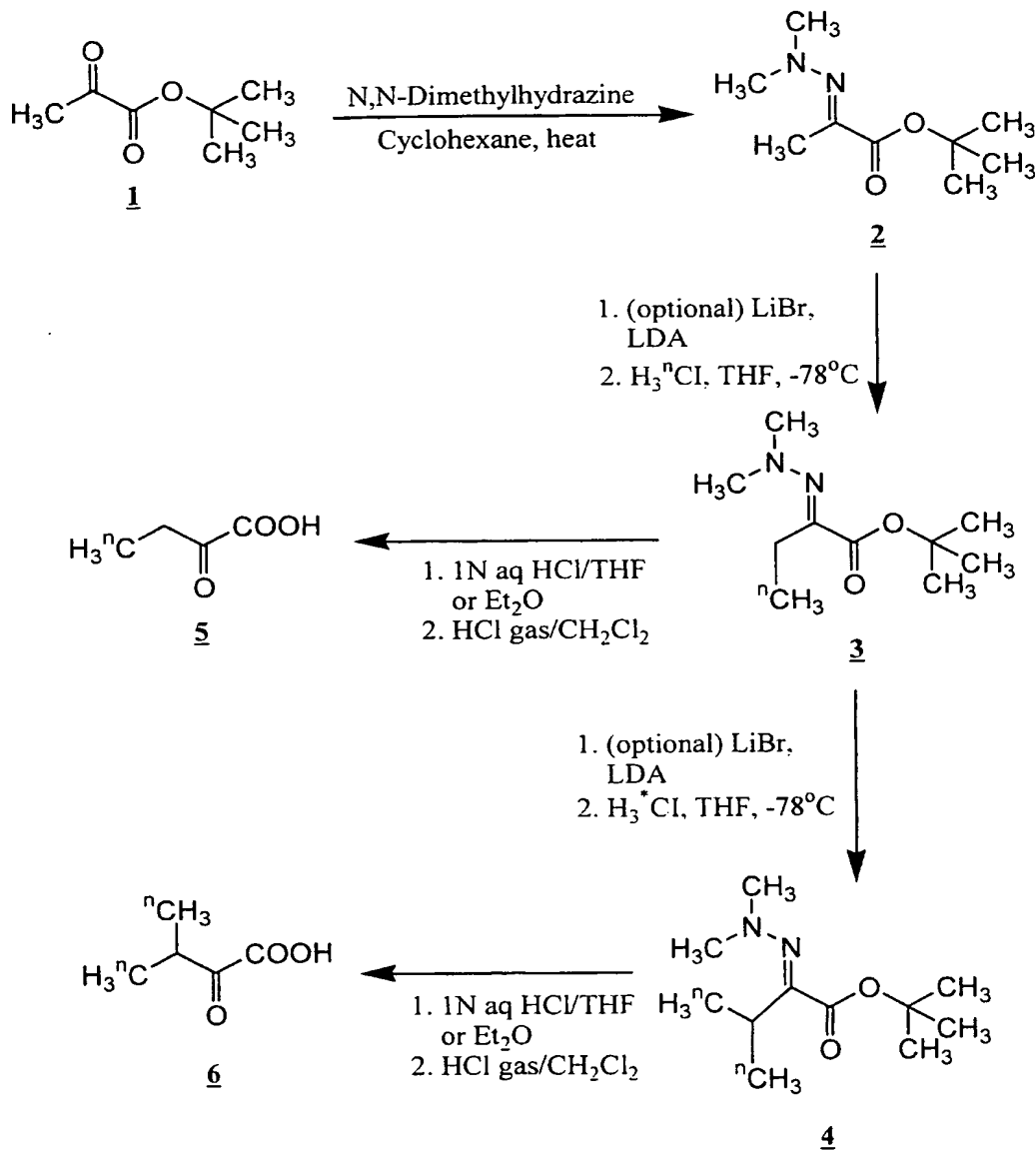
Reaction Schemes II, III, and IV illustrate, respectively, how these α-ketoacids are biosynthetically converted into ^{13}C-leucine, ^{13}C-isoleucine, and ^{13}C-valine. In all of the Schemes, the sites of isotopic enrichment are indicated by asterisks.

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Scheme II

Biochemical Synthesis of Labeled Isoleucine

H_3^*C —COOH (**5**) →[Acetolactate Synthase; Pyruvate → CO_2]→ (**7**)

(**7**) →[Ketol-Acid Reductoisomerase; NADH → NAD^+]→ (**8**)

(**8**) →[1. Dihydroxy acid Dehydratase 2. Reductase; → H_2O]→ (**9**)

(**9**) →[Branched-Chain Acid Transaminase; L-Glutamate → 2-Oxoglutamate]→ (**10**) Isoleucine

5

10

15
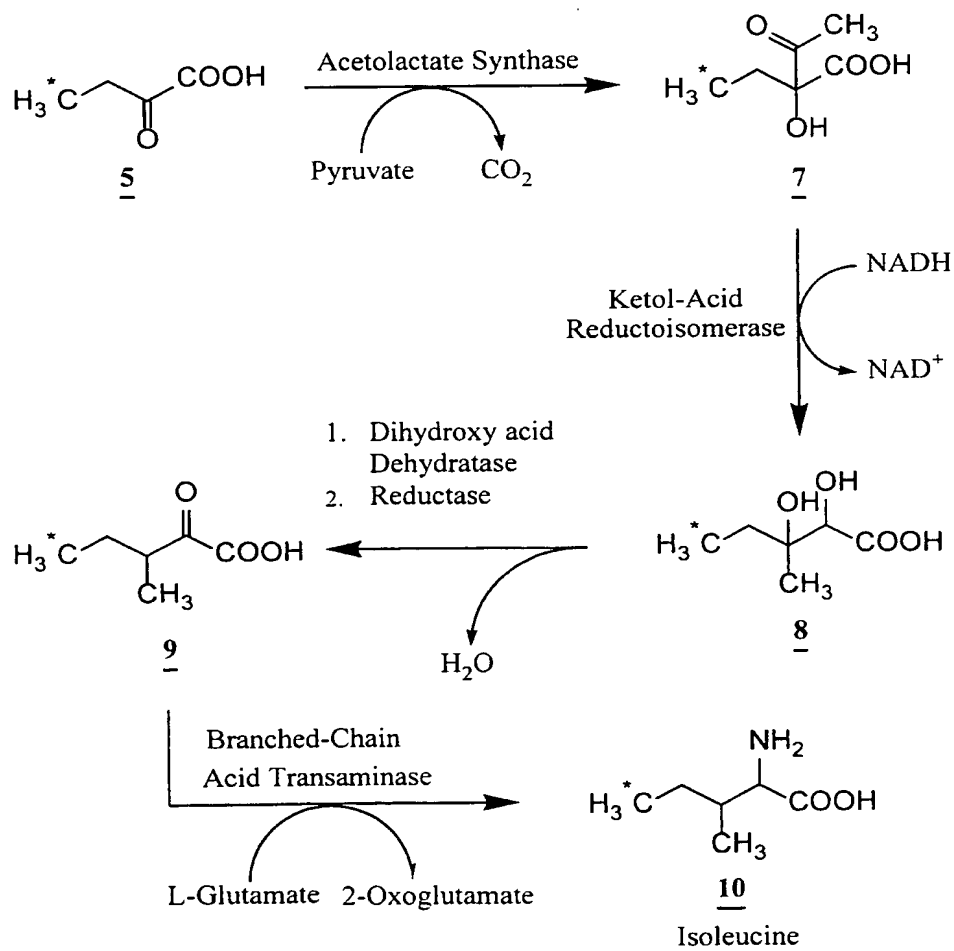
Scheme III

Biochemical Synthesis of Labeled Leucine

6

2-Isopropylmalate Synthase

Acetyl-CoA
H₂O

CoA

11

3-Isopropylmalate
Dehydratase

H₂O

3-Isopropylmalate
Dehydratase

12

H₂O

13

NAD⁺

3-Isopropylmalate
Dehydrogenase

NADH

CO₂

15

Leucine
Transaminase

L-Glutamate

2-Oxo-glutamate

14

16
Leucine

17
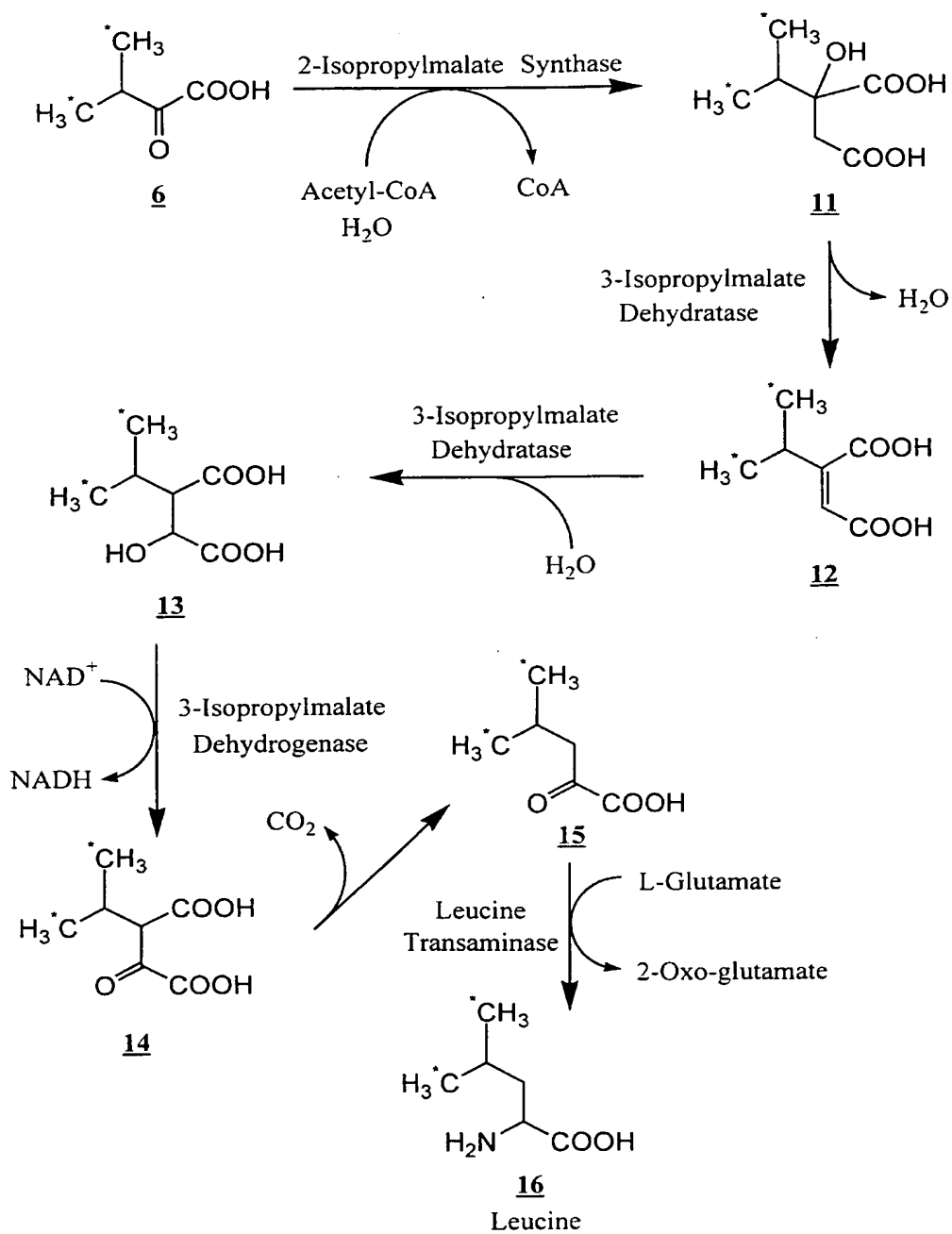
Scheme IV

Biochemical Synthesis of Valine

6   L-Alanine   Pyruvate   **17**

                                                        L-Valine
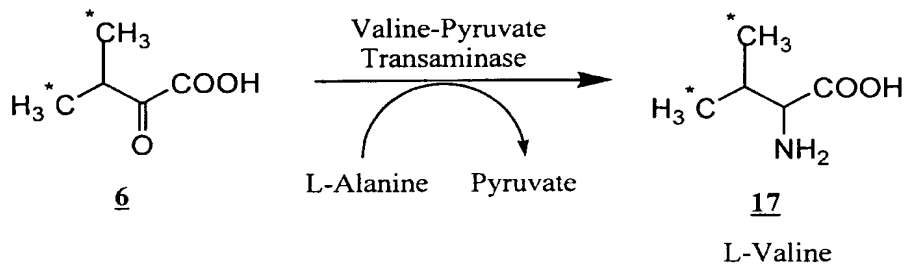
Means for preparing expression vectors that contain
polynucleotide sequences coding specific polypeptides and for
transforming host cells with those vectors are well known in
the art. (*See*, for example, R. W. Old, *et al.*, "Techniques of
Gene Manipulation," Blackwell Science, London, 1994, and
similar treatises in the field.) Likewise, methods for
culturing the transformed cells to express the coded
polypeptide and for isolating, purifying and re-folding the
polypeptide are also well known in the art. Examples
presented below describe the production from modified *E. coli*
of ^{13}C-enriched samples of the 81-256 amino acid catalytic
region of human stromelysin and FK binding protein (FKBP), and
the use of these isotopically-enriched polypeptides in the
instant methods.

Where the method is employed to screen more than one
compound for binding to the target molecule, for example a
mixture or a library of compounds, and where a difference
arises between the first spectrum generated from the target
molecule alone and that generated from the target molecule in

the presence of compound(s), additional steps are performed to
identify which specific compound or compounds contained in the
mixture is actually binding to the target molecule. Those
additional steps include exposing the ^{13}C-enriched target
5   molecule individually to each compound of the mixture;
generating a two-dimensional ^{13}C/^{1}H NMR correlation spectrum of
the labeled target molecule that has been individually exposed
to each compound; and comparing each spectrum to the first
spectrum generated from the target molecule alone to determine
10   differences in any of those compared spectra. The differences
in the spectra facilitate the identification of a compound
that is a ligand.

The chemical shift values of the particular ^{13}C/^{1}H signals
in the two dimensional correlation spectrum correspond to
15   known specific locations of atomic groupings in the target
molecule (for example, the carbon atoms of a particular amino
acid residue in the target molecule or, in the case of a
polypeptide specifically labeled at the methyl groups of
alanyl, leucyl, isoleucyl, and valinyl residues). The
20   screening process of this invention thus allows not only for
the identification of which compound(s) bind to a particular
target molecule, but also permits the determination of the
particular amino acids that are affected by the binding of the
compound to the target molecule. The chemical shift values
25   may reflect a change in the conformation of the target
molecule, or may reflect the binding of the ligand compound at
the site that corresponds to that particular signal.

In addition, the dissociation constant, K_D, for a given
ligand and its target molecule can be determined by this

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process, if desired, by generating a first two-dimensional ^{13}C /^{1}H NMR correlation spectrum of a ^{13}C-labeled target molecule; exposing the labeled target molecule to various concentrations of a ligand; generating a two-dimensional ^{13}C/^{1}H NMR

5   correlation spectrum at each concentration of ligand employed; comparing the spectra generated to the first spectrum of the target molecule; and calculating the dissociation constant between the target molecule and the ligand from those differences using Equation 1:

10

$$K_D = \frac{([P_0] - x)([L_0] - x)}{x}$$

where [P_0] is the total molar concentration of the target molecule, [L_0] is the total molar concentration of the ligand,

15   and x is the molar concentration of the bound species. The value of x is determined from the NMR chemical shift data according to Equation 2:

$$x = \frac{\delta_{observed} - \delta_{free}}{\Delta}$$

20

where $\delta_{observed}$ is the observed chemical shift value, δ_{free} is the chemical shift value for the free species, and Δ is the difference between the limiting chemical shift value for saturation ($\delta_{saturation}$) and the chemical shift value of the target

25   molecule free of bound ligand (δ_{free}).

The dissociation constant is then determined by varying its value until a best fit is obtained with the observed data

using standard curve-fitting statistical methods. In those
situations where the value of $\delta_{saturation}$ is not directly known, K_D
and $\delta_{saturation}$ are varied and the resulting data subjected to the
same curve-fitting statistical method.

5   An advantageous capability of the screening method is its
ability to determine the dissociation constant of one ligand
of the target molecule in the presence of a second molecule
already bound to the ligand. This is generally not possible
with other methods which employ "wet chemical" analytical
10   methods of determining binding of a ligand to a target
molecule substrate.

       The process of determining the dissociation constant of a
ligand can be performed in the presence of a second bound
ligand. Accordingly, the ^{13}C-labeled target molecule is bound
15   to that second ligand before exposing that target to the test
compounds. The screening method is additionally able to
provide information regarding the binding of a second or
subsequent ligand to the target molecule. This second ligand
may be chemically linked to the first ligand bound to the
20   target molecule, thus providing a new composite molecule for
use in affecting the target molecule.

       The screening method of the present invention begins with
the generation or acquisition of a two-dimensional $^{13}C/^{1}H$
correlation spectrum of the isotopically enriched target
25   molecule. As stated above, the target molecule can be either
uniformly enriched with ^{13}C, or it can be specifically enriched
by the incorporation of ^{13}C-methyl groups in alanyl, leucyl,
valinyl, and isoleucyl residues. Means for generating two-
dimensional $^{13}C/^{1}H$ correlation spectra are well known in the

art. The NMR spectra that are typically recorded are two-
dimensional $^{13}C/^1H$ heteronuclear single quantum correlation
(HSQC) spectra, although other techniques known to those
skilled in the art can be used. Because the $^{13}C/^1H$ signals

5   corresponding to the protein are usually well resolved, the
chemical shift changes for individual $^{13}C/^1H$ pairs can be
readily monitored. A representative two-dimensional $^{13}C/^1H$
correlation spectrum of a ^{13}C-labeled target polypeptide (FKBP)
is shown in Figure 1. A representative two-dimensional $^{13}C/^1H$

10   correlation spectrum of a specifically ^{13}C-enriched FKBP is
shown in Figure 3.

        Following exposure of the ^{13}C-labeled target molecule to
one or more test compounds, a second two-dimensional $^{13}C/^1H$ NMR
correlation spectrum is generated. That spectrum is generated

15   in the same manner as set forth above. The first and second
spectra are then compared to determine whether there are any
differences between the two spectra. Differences in the two-
dimensional $^{13}C/^1H$ NMR correlation spectra that indicate the
presence of a ligand correspond to ^{13}C-labeled sites in the

20   target molecule. Those differences are determined using
standard procedures well known in the art. Figure 2 shows the
methyl regions of uniformly ^{13}C-labeled FKBP before (thin
multiple contours) and after (thick single contours) addition
of 2-phenylimidazole (0.12 mM).

25   Particular signals in a two-dimensional $^{13}C/^1H$ correlation
spectrum correspond to specific carbon and proton atoms in the
target molecule (for example, particular methyl groups of the
amino acid residues in the protein). By way of example, it
can be seen from Figure 3 that chemical shifts observed in

two-dimensional $^{13}C/^{1}H$ correlation spectra of FKBP exposed to a
test compound occurred at residue positions 97 (leucine) and
55 (valine). The region of the protein that is responsible
for binding to the individual compounds is identified from the
particular carbon and proton atom pairs that change upon the
addition of compound.

EXAMPLES

Preparation 1

5   **Preparation of Uniformly ^{13}C-Enriched Catalytic Domain of Human Stromelysin (SCD)**

The 81-256 fragment (SEQ ID NO: 1) of stromelysin (SCD) is prepared by inserting a plasmid which codes for the

10   production of the protein fragment into an *E. coli* strain and growing the genetically-modified bacterial strain in a suitable culture medium. The protein fragment is isolated from the culture medium, purified, and subsequently used in the two-dimensional NMR analysis of its affinity with test

15   compounds in accordance with the method of this invention. The procedures for the preparation processes are described below.

Human skin fibroblasts (ATCC No. CRL 1507) are grown and induced using the procedure described by Clark *et al.*, *Archiv.*

20   *Biochem. and Biophys.* 241: 36 (1985). Total RNA is isolated from 1 g of cells using a RNAgents® Total RNA Isolation System Kit (Promega Corp.) following the manufacturer's instructions. A 1 μg portion of the RNA is denatured by heating at 80°C for five minutes and then subjected to reverse transcriptase PCR

25   using a GenAmp® RNA PCR kit (Applied Biosystems/Perkin-Elmer) following the manufacturer's instructions.

Nested PCR is performed using first primers (a) GAAATGAAGAGTCTTCAA (SEQ ID NO: 2) and (b) GCGTCCCAGGTTCTGGAG (SEQ ID NO. 3) and thirty-five cycles of 94°C, two minutes; 45°

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C, two minutes; and 72°C, three minutes. This is followed by
re-amplification with internal primers (c)
TACCATGGCCTATCCATTGGATGGAGC (SEQ ID NO: 4) and (d)
ATAGGATCCTTAGGTCTCAGGGGA GTCAGG (SEQ ID NO: 5) using thirty

5   cycles under the same conditions described immediately above
to generate a DNA sequence coding for amino acid residues
1-256 of human stromelysin.

        The PCR fragment is then cloned into PCR cloning vector
pT7BIue® (Novagen, Inc.) according to the manufacturer's

10   instructions. The resulting plasmid is cut with NcoI and
BamHI and the stromelysin fragment is sub-cloned into the
expression vector pET3d (Novagen, Inc.).

        A mature stromelysin expression construct coding for
amino acid residues 81-256 plus an initiating methionine

15   aminoacyl residue is generated from the 1-256 expression
construct by PCR amplification. The resulting PCR fragment is
first cloned into the pT7BIue® vector (Novagen, Inc.) and then
sub-cloned into the pET3d vector (Novagen, Inc.), using the
manufacturer's instructions in the manner described above, to

20   produce plasmid pETST-83-256. This final plasmid is identical
to that described by Qi-Zhuang et al., Biochemistry, 31: 11231
(1992) with the exception that the present plasmid codes for a
peptide sequence beginning two amino acids earlier, at
position 81, in the sequence of human stromelysin. Plasmid

25   pETST-83-256 is transformed into E. coli strain
BL21(DE3)/pLysS (Novagen, Inc.) in accordance with the
manufacturer's instructions to generate an expression strain,
BL21(DE3)/pLysS/pETST-255-1.

A pre-culture medium is prepared by dissolving 1.698 g of $NaH_2PO_4 \bullet 7H_2O$, 0.45 g of KH_2PO_4, 0.075 g NaCl, 0.150 g NH_4Cl, 0.3 g U-^{13}C-glucose, 300 μl of 1M aqueous $MgSO_4$ solution, and 15 mL of aqueous $CaCl_2$ solution in 150 ml of deionized water. The

5   resulting solution of pre-culture medium is sterilized and transferred to a sterile 500 ml baffle flask. Immediately prior to inoculation of the pre-culture medium with the bacterial strain, 150 ml of a solution containing 34 mg/ml, of chloramphenicol in 100% ethanol and 1.5 ml of a solution

10   containing 20 mg/ml of ampicillin is added to the flask contents. The flask contents are then inoculated with 1 ml of glycerol stock of genetically modified *E. coli* strain BL21(DE3)/pLysS/pETST-255-1. The flask contents are shaken (225 rpm) at 37°C until an optical density of 0.65 is observed.

15   A fermentation nutrient medium is prepared by dissolving 113.28 g of $Na_2HPO_4 \bullet 7H2O$, 30 g of KH_2PO_4, 5 g NaCl and 10 mL of 1% DF-60 antifoam agent in 9604 mL of deionized water. This solution is placed in a New Brunswick Scientific Micros Fermenter and sterilized at 121 °C for 40 minutes.

20   Immediately prior to inoculation of the fermentation medium, the following pre-sterilized components are added to the fermentation vessel contents: 100 ml of a 10% aqueous solution of NH_4Cl, 15 g of uniformly ^{13}C-enriched glucose, 20 ml of an aqueous 1M solution of $MgSO_4$, 1 ml of an aqueous 1M $CaCl_2$

25   solution, 5 ml of an aqueous solution of thiamin hydrochloride (10 mg/ml), 10 ml of a solution containing 34 mg/ml of chloramphenicol in 100% ethanol, and 1.9 g of ampicillin dissolved in the chloramphcnicol solution. The pH of the

resulting solution is adjusted to pH 7.00 by the addition of an aqueous solution of 4N H_2SO_4.

The pre-culture of *E. coli* strain BL21(DE3)/pLysS/pETST-255-1 from the shake flask scale

5   procedure described above is added to the fermenter contents and cell growth is allowed to proceed until an optical density of 0.48 is achieved. During this process, the fermenter contents are automatically maintained at pH 7.0 by the addition of 4N H_2SO_4 or 4N KOH as needed. The dissolved oxygen

10   content of the fermenter contents is maintained above 55% air saturation through a cascaded loop which increased agitation speed when the dissolved oxygen content dropped below 55%. Air is fed to the fermenter contents at 7 standard liters per minute (SLPM) and the culture temperature is maintained at

15   37°C throughout the process.

The cells are harvested by centrifugation at 17,000 x g᠄ for 10 minutes at 4°C and the resulting cell pellets are collected and stored at -85°C. The wet cell yield is 3.5 g/L. Analysis of the soluble and insoluble fractions of cell

20   lysates by sodium dodecyl sulfate polyacrylamide gel electrophoresis (SDS-PAGE) reveals that approximately 50% of the stromelysin is found in the soluble phase.

The stromelysin fragment prepared as described above is purified employing a modification of the technique described

25   by Ye, *et al.*, *Biochemistry*, 31: 11231 (1992). The harvested cells are suspended in 20 mM Tris-HCl buffer (pH 8.0), sodium azide solution containing 1mM $MgCl_2$, 0.5 mM $ZnCl_2$, 25 units/ml of Benzonase® enzyme (Benzon Pharma), and an inhibitor mixture made up of 4-(2-aminoethyl)benzenesulfonyl fluoride ("AEBSF")

Leupeptin®, Aprotinin® and Pepstatin® (all at concentrations of 1 mg/ml). AEBSF, Leupeptin®, Aprotinin®, and Pepstatin® are available from American International Chemical. The resulting mixture is gently stirred for one hour and then

5 cooled to 4°C. The cells are then sonically disrupted using a 50% duty cycle. The resulting lysate is centrifuged at 14,000 rpm for 30 minutes and the pellet of insoluble fraction frozen at -80°C for subsequent processing.

Solid ammonium sulfate is added to the supernatant to the

10 point of 20% of saturation and the resulting solution loaded onto a 700 ml phenyl Sepharose fast flow ("Q-Sepharose FF) column (Pharmacia Biotech.). Prior to loading, the Sepharose column is equilibrated with 50 mM Tris-HCl buffer (pH 7.6 at 4°C), 5 mM CaCl2, and 1M $(NH_4)_2SO_4$. The loaded column is

15 eluted with a linear gradient of decreasing concentrations of aqueous $(NH_4)_2SO_4$ (from 1M down to 0M) and increasing concentrations of aqueous $CaCl_2$ (from 5 mM to 20 mM) in Tris-HCl buffer at pH 7.6. The active fractions of eluate are collected and concentrated in an Amicon stirred cell (Amicon,

20 Inc.). The concentrated sample is dialyzed overnight in the starting buffer used with the Q-Sepharose FF column, 50 mM Tris-HCl (pH 8.2 at 4°C) with 10 mM $CaCl_2$.

The dialyzed sample is then loaded on the Q-Sepharose FF column and eluted with a linear gradient comprising the

25 starting buffer and 200 nM NaCl. The purified soluble fraction of the stromelysin fragment is concentrated and stored at 4°C. The pellet is solubilized in 8M guanidine-HCl. The solution is centrifuged for 20 minutes at 20,000 rpm and the supernatant added dropwise to a folding buffer comprising

50 mM Tris-HCl (pH 7.6), 10 mM $CaCl_2$, 0.5 mM $ZnCl_2$ and the inhibitor cocktail of AEBSF, Leupeptin®, Aprotinin®, and Pepstatin® (all at concentrations of 1 µg/ml). The volume of folding buffer is ten times that of the supernatant. The

5   mixture of supernatant and folding buffer are centrifuged at 20,000 rpm for 30 minutes. The supernatant from this centrifugation is stored at 4°C and the pellet subjected twice to the steps described above of solubilization in guanidine-HCl, refolding in buffer, and centrifugation. The

10   final supernatants from each of the three centrifugations are combined and solid ammonium sulfate was added to the point of 20% saturation. The resulting solution thus derived from the insoluble fraction is subjected to purification on phenyl Sepharose and Q-Sepharose as described above for the soluble

15   fraction. The purified soluble and insoluble fractions are combined to produce about 1.8 mg of purified stromelysin 81-256 fragment (SCD) per gram of original cell paste, uniformly enriched with ^{13}C.

20

Preparation 2

Preparation of Specifically ^{13}C-Enriched Catalytic Domain of Human Stromelysin (SCD)

25   SCD is expressed by culturing the BL21(DE3)/pLysS/pETST-255-1 modified *E. coli* strain in a medium comprising ^{13}C-enriched α-ketobutyric acid and α-keto-isovaleric acid. The methods used for preparation of the genetically-engineered strain of *E. coli*, and for expressing,

isolating, and purifying the protein fragment are as described above, except for the use of $U-^{12}C$-glucose, instead of $U-^{13}C$-glucose.

<div align="center">

Preparation 3

Preparation of Uniformly and Specifically ^{13}C-Enriched FKBP

</div>

A. Preparation of Specifically ^{13}C-Enriched Val/Leu FKBP

Cells transformed with a plasmid encoding for human FKBP (as described in Egan, *et al.*, *Biochemistry* 32: 1920-1927 (1993)) were grown in a 100% D_2O culture medium containing $^{15}NH_4Cl$ (Cambridge Isotopes) as the sole nitrogen source (1.0 g/L) and perdeuterated glucose (1.5 g/L) as the carbon source (Cambridge Isotopes). Cells were grown at 37°C and the flask contents were shaken until an optical density of 1.0 was obtained. One hour prior to induction, 80 mg of L-valine-U-$^{13}C_5-^{15}N-2,3-d_2$ (Cambridge Isotopes) was added to the culture medium. The culture was induced with 1 mM IPTG for 12 hours.

The cells were harvested by centrifugation at 17,000 x g for 10 minutes at 4 °C and the resulting cell pellets were suspended in 50 mM phosphate buffer (pH 7.4) containing 5 mM DTT and 1 mM PMSF and mechanically lysed using a French press. The resulting lysate was centrifuged at 25,000 rpm for 30 minutes. Solid ammoniuum sulfate was added to the supernatant to the point of 40% of saturation and the centrifuged at 18,000 rpm for 5 minutes. The supernatant was dialyzed into 10 mM HEPES (pH 8.0) for 12 hours at 4 °C. The resulting

solution was then loaded onto a 10 mL Q-Sepharose fast flow
column (Sigma) pre-equilibrated in the dialysis buffer.
Fractions were collected, pooled, and concentrated using an
Amicon flow cell. The solution was then dialyzed into 20 mM

5   phosphate buffer (pH 6.5) containing 10 mM DTT and 0.01%
sodium azide.

B. Preparation of Specifically ^{13}C-Enriched
Val/Leu/Ile FKBP

10

Cells transformed with a plasmid encoding for human FKBP
(as described in Egan, *et al.*, *Biochemistry* 32: 1920-1927
(1993)) are grown in a culture medium containing NH$_4$Cl as the
sole nitrogen source (1.0 g/L) and glucose (1.5 g/L) as the

15   carbon source. Cells are grown at 37°C and the flask contents
are shaken until an optical density of 1.0 is obtained. One
hour prior to induction, 100 mg of α-keto-butyric acid and α-
keto-isovaleric acid are added to the culture medium. The
culture is induced with 1 mM IPTG for 12 hours. Expression,

20   isolation, and purification of the expressed protein are as
described above.

C. Preparation of Uniformly ^{13}C-Enriched FKBP

25   Cells transformed with a plasmid encoding for human FKBP
(as described in Egan, *et al.*, *Biochemistry* 32: 1920-1927
(1993)) were grown in a culture medium containing ^{15}NH$_4$Cl
(Cambridge Isotopes) as the sole nitrogen source (1.0 g/L) and
uniformly ^{13}C-enriched glucose (1.5 g/L) as the carbon source

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(Cambridge Isotopes). Expression, isolation, and purification
of the expressed protein are as described above.

5

Example 1
Screening for Ligands Using $^{13}C/^{1}H$ NMR Correlation Spectra Using Uniformly ^{13}C-Enriched FKBP

10   Uniformly ^{13}C-enriched FKBP was prepared in accordance
with the procedures detailed above. The protein solutions
used in the screening assay contained the uniformly ^{13}C-
enriched FKBP (0.2 mM) and sodium azide (0.05%) in an H_2O/D_2O
(9/1) phosphate buffered solution (20 mM, pH 6.5).

15   Two-dimensional $^{13}C/^{1}H$ NMR spectra were generated at 30°C
on a Bruker DRX500 NMR spectrometer equipped with a triple
resonance probe and Bruker sample changer. The $^{13}C/^{1}H$ HSQC
spectra were acquired as 64 x 1024 complex points using sweep
widths of 3771 Hz (^{13}C, t_1) and 8333 Hz (^{1}H, t_2). A delay of 1
20   second between scans and 8 scans per free induction decay
(fid) were employed in the data collection. All NMR spectra
were processed and analyzed on Silicon Graphics computers.

       A first two-dimensional $^{13}C/^{1}H$ NMR correlation spectrum
was acquired for the ^{13}C-labeled FKBP target molecule as
25   described above. The FKBP target was then exposed to a
library mixture of test compounds. Stock solutions of the
compounds were made at 100 mM and 1 M. In addition, a
combination library was prepared which contained 8-10

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compounds per sample at a concentration of 100 mM for each compound.

The pH of the 1 M stock solution was adjusted with acetic acid and ethanolamine so that no pH change was observed upon a
5 1/10 dilution with a 100 mM phosphate buffered solution (pH 7.0). It is important to adjust the pH, because small changes in pH can alter the chemical shifts of the biomolecules and complicate the interpretation of the NMR data.

The compounds in the library were selected on the basis
10 of size (molecular weight = 100-300) and molecular diversity. The molecules in the collection had different shapes (for example, flat aromatic rings(s), puckered aliphatic rings(s), straight and branched chain aliphatics with single, double, or triple bonds) and diverse functional groups (for example,
15 carboxylic acids, esters, ethers, amines, aldehydes, ketones, and various heterocyclic rings) to maximize the possibility of discovering compounds that interact with widely diverse binding sites.

The NMR samples were prepared by adding 1.25 μl of the
20 dimethyl sulfoxide stock solution of the compound mixtures that contained each compound at a concentration of 100 mM to 0.5 ml H_2O/D_2O (9/1) buffered solution of the uniformly ^{13}C-labeled protein. The final concentration of each of the compounds in the NMR sample was about 0.25 mM.
25 In the screening experiment, one compound, 2-phenylimidazole, was found to bind to FKBP. Figure 1 shows the $^{13}C/^1H$ correlation spectrum of uniformly ^{13}C-labeled FKBP. The spectrum (128 complex points, 4 scans/fid) was acquired on a

33

0.1 mM sample of FKBP in 20 mM phosphate (pH 6.5), 0.01%
sodium azide and 10% deuterium oxide (D_2O).

Figure 1 shows a $^{13}C/^1H$ correlation spectrum of uniformly
^{13}C-labeled FKBP. The spectrum (128 complex points, 4

5   scans/fid) was acquired on a 0.1 mM sample of FKBP in 20 mM
phosphate (pH 6.5), 0.01% sodium azide and 10% deuterium oxide
(D_2O). Figure 2 shows the methyl regions of $^{13}C/^1H$ correlation
spectra (64 complex points, 8 scans/fid) of uniformly ^{13}C-
labeled FKBP (0.2 mM) before (thin multiple contours) and

10   after (thick single contours) the addition of 2-
phenylimidazole (0.25 mM). The changes in chemical shifts at
aminoacyl residues Leu[97], Val[55], Ile[56] and Ile[90] are
indicated.

Figure 3 shows $^{13}C/^1H$ correlation spectra (48 complex

15   points, 8 scans/fid) of FKBP selectively $^{13}C/^{15}N/^2H$ labeled at
valinyl and leucyl residues before (thin multiple contours)
and after (thick single contours) addition of 2-
phenylimidazole (0.25 mM). All other conditions are the same
as those employed in generating the spectra illustrated in

20   Figure 1. Selected residues that show significant changes
upon binding are indicated. Again, changes in the chemical
shift values indicate that binding is occurring at or near the
Leu[97] and Val[55] residues.

Figure 4 shows $^{13}C/^1H$ correlation spectra of FKBP

25   selectively ^{13}C labeled at alanyl residues before (thin
multiple contours) and after (thick single contours) addition
of 2-phenylimidazole (0.25 mM). The super-position of the
chemical shift values before (light multiple contours) and

34

after (heavy single contour) addition of the ligand indicate that none of the alanyl residues are involved in the binding.

Figure 5 shows a "stick model" depiction of the three-dimensional structure of FKBP. Selected residues are numbered for aid in visualization. The aminoacyl residues involved in the binding site in the protein have been shown in bold.

35

WE CLAIM:

    1.   A method of detecting binding between a putative
ligand and a pre-selected, ^{13}C-enriched target molecule
which comprises:

5   a)   generating a first two-dimensional ^{13}C/^{1}H NMR
correlation spectrum of said target molecule;

         b)   forming a mixture of said target molecule with at
least one putative ligand compound;

10

         c)   generating a second two-dimensional ^{13}C/^{1}H NMR
correlation spectrum of the mixture of step (b); and

         d)   comparing the first and second spectra.

15

    2.   The method of Claim 1 wherein said target
molecule is selected from the group consisting of
lipoproteins, lipoprotein fragments, glycoproteins,
glycoprotein fragments, proteins, protein fragments,
5 polypeptides, DNA, and RNA.

    3.   The method of Claim 1 wherein said target
molecule is selected from the group consisting of proteins,
protein fragments, and polypeptides.

    4.   The method of Claim 3 wherein said target
molecule is prepared by culturing a transformed cell line

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which contains an expression vector containing a polynucleotide encoding said target molecule in a medium
5   containing assimilable sources of ^{13}C.

5.   The method of Claim 4 wherein said assimilable sources of ^{13}C are uniformly ^{13}C-enriched.

6.   The method of Claim 5 wherein uniformly ^{13}C-labeled glucose (U-^{13}C-glucose) is employed to produce said assimilable sources of ^{13}C.

7.   The method of Claim 4 wherein said assimilable sources of ^{13}C are specifically ^{13}C-enriched.

8.   The method of Claim 7 wherein ^{13}C-enriched methyl iodide is employed to produce said assimilable sources of ^{13}C.

9.   The method of Claim 7 wherein said assimilable sources of ^{13}C are amino acids, and salts thereof.

10.   The method of Claim 9 wherein said amino acid is selected from the group consisting of alanine, leucine, isoleucine, and valine.

11.   The method of Claim 8 wherein said assimilable sources of ^{13}C are biosynthetic precursors of amino acids, and salts thereof.

12. The method of Claim 11 wherein said biosynthetic precursors of amino acids are selected from the group consisting of $4(^{13}C)$-butyric acid and $4-(^{13}C)-3-(^{13}C)$-methylbutyric acid.

5

13. A method of screening a mixture of compounds for binding to a pre-selected, ^{13}C-enriched target molecule which comprises:

5   a)   generating a first two-dimensional $^{13}C/^1H$ NMR correlation spectrum of said target molecule;

        b)   contacting said target molecule with said mixture of compounds;

10

        c)   generating a second two-dimensional $^{13}C/^1H$ NMR correlation spectrum of the mixture of step (b);

        d)   comparing the first and second spectra.

15

14. The method of Claim 13, wherein said method additionally comprises:

5   e)   exposing said target molecule individually to each compound in said mixture when step d) reveals differences in the first and second spectra;

f)   generating two-dimensional $^{13}C/^1H$ NMR correlation
10   spectra of said target molecule that has been exposed to
each compound; and

g)   comparing each spectra generated in step f) to
the first spectrum generated from the target molecule
15   alone.

15.   The method of Claim 14 wherein said target
molecule is selected from the group consisting of
lipoproteins, lipoprotein fragments, glycoproteins,
glycoprotein fragments, proteins, protein fragments,
5   polypeptides, DNA, and RNA.

16.   The method of Claim 15 wherein said target
molecule is selected from proteins, protein fragments, and
polypeptides.

17.   The method of Claim 13 wherein said target
molecule is uniformly ^{13}C-enriched.

18.   The method of Claim 13 wherein said target
molecule is specifically ^{13}C-enriched.

19.   A method of determining the dissociation constant
for a ligand which binds to a pre-selected, ^{13}C-enriched
target molecule which comprises:

5   a)   generating a first two-dimensional ^{13}C/^{1}H NMR
correlation spectrum of said target molecule;

           b)   exposing said target molecule to various
concentrations of said ligand;

10

           c)   generating a two-dimensional ^{13}C/^{1}H NMR
correlation spectrum at each concentration of ligand in
step b);

15   d)   comparing each spectrum from step (c) to said
first spectrum from step (a); and

           e)   calculating the dissociation constant.

           20.   The method of Claim 19 wherein said target
molecule is selected from the group consisting of
lipoproteins, lipoprotein fragments, glycoproteins,
glycoprotein fragments, proteins, protein fragments,
5   polypeptides, DNA, and RNA.

           21.   The method of Claim 20 wherein said target
molecule is selected from proteins, protein fragments, and
polypeptides.

           22.   The method of Claim 20 wherein said target
molecule is uniformly ^{13}C-enriched.

40

23. The method of Claim 20 wherein said target
molecule is uniformly ^{13}C-enriched.

24. A compound identified by the screening method of
Claim 13.

25. A method of determining the specific amino acid
residues in a pre-selected, ^{13}C-enriched target molecule
affected by the binding of a ligand to said target molecule
which comprises:

5

a)   generating a first two-dimensional ^{13}C/^{1}H NMR
correlation spectrum of said target molecule, wherein said
chemical shift values of the ^{13}C/^{1}H signals in said two
dimensional correlation spectrum correspond to at least one
10   known specific location of atomic groupings in said target
molecule;

b)   forming a mixture of said target molecule with a
known ligand compound;

15

c)   generating a second two-dimensional ^{13}C/^{1}H NMR
correlation spectrum of the mixture of step (b); and
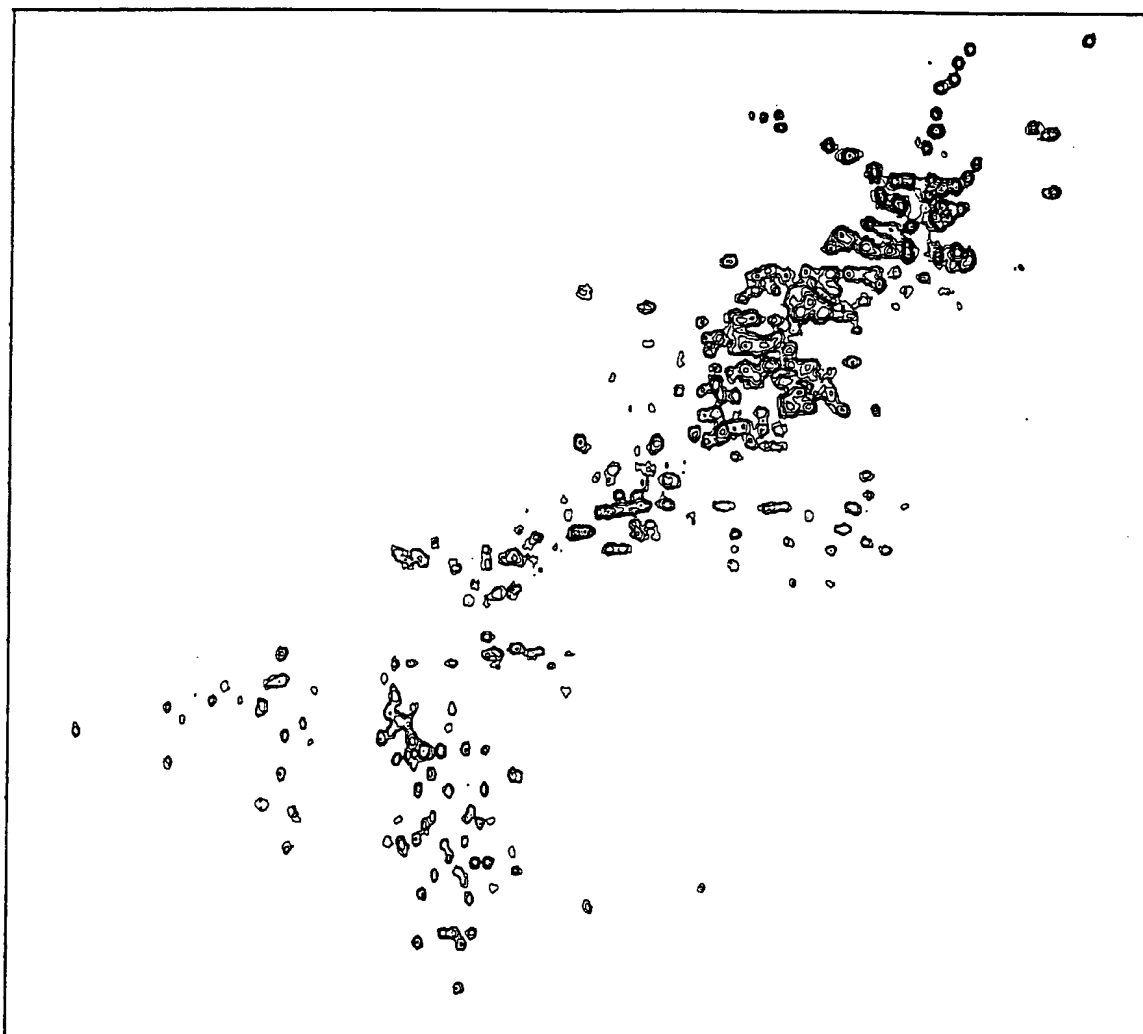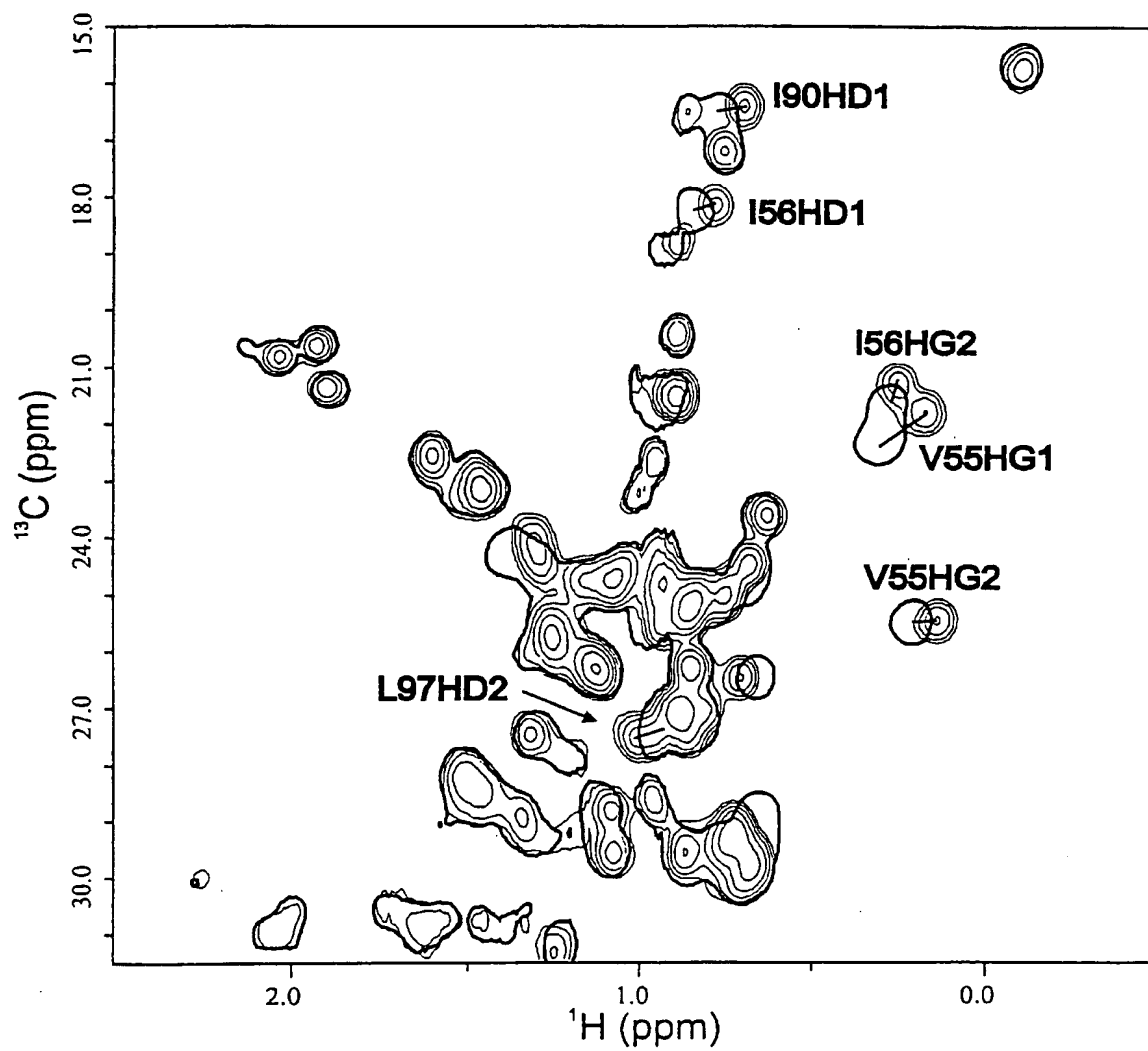
d)   comparing the first and second spectra.

20

FIG.1

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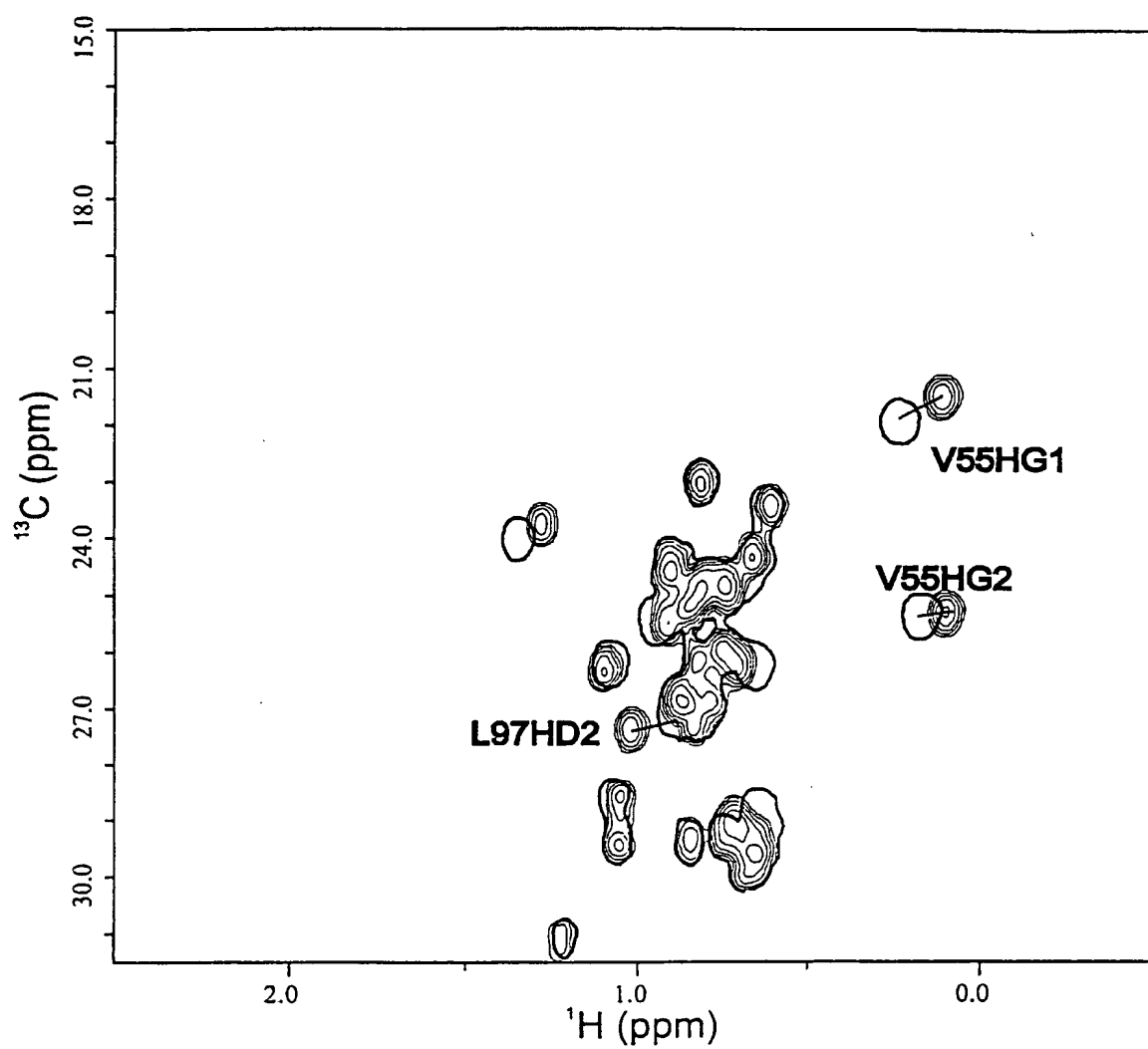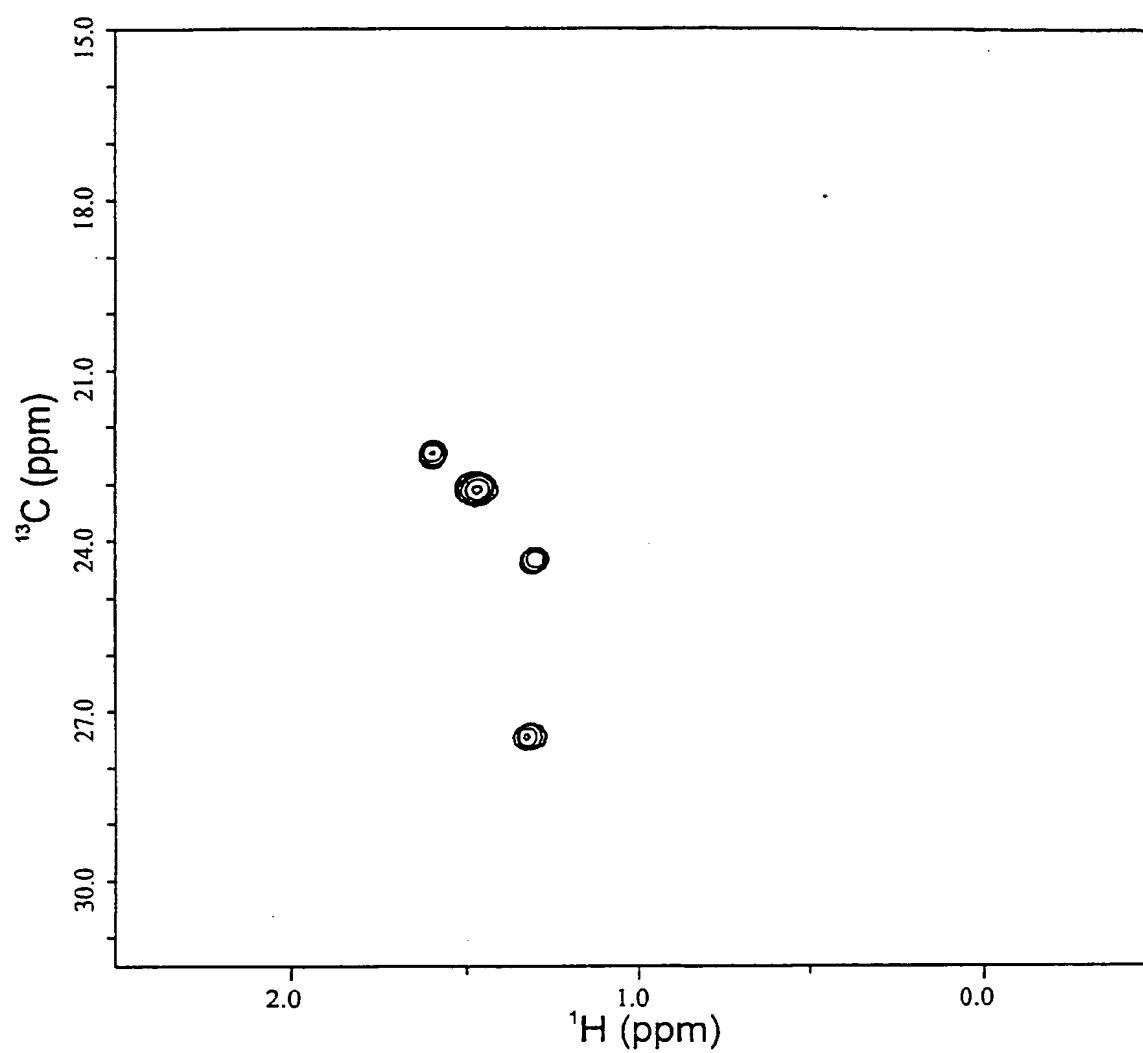
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FIG.2

FIG.3

FIG.4

5/5

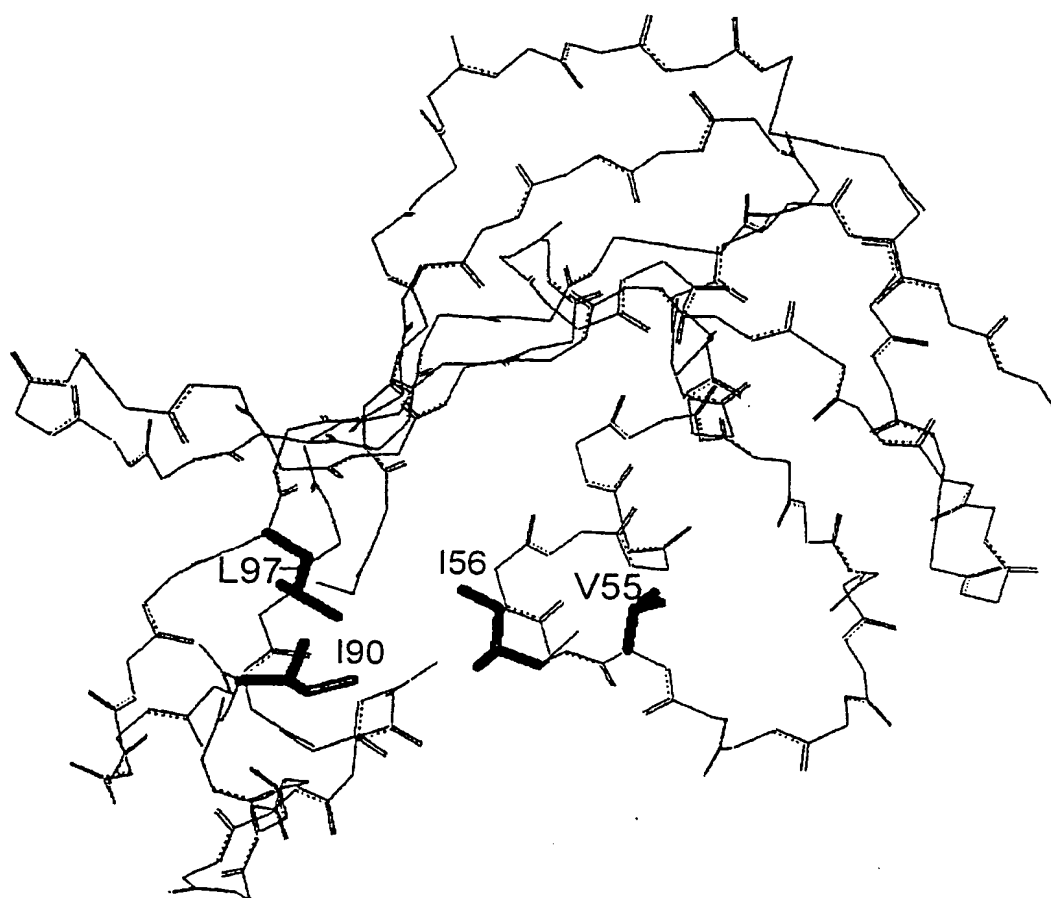

FIG.5

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SEQUENCE LISTING

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      Hajduk, P. J.

<120> USE OF 13C NUCLEAR MAGNETIC RESONANCE TO
      DETECT BINDING TO TARGET BIOMOLECULES

<130> 5816.US.P3

<140> US 09/288,924
<141> 1999-04-09

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<151> 1999-02-01

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INTERNATIONAL SEARCH REPORT

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7   G01N33/68   G01N33/94   G01N33/58   C07B61/00   G01N33/50

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7   G01N   C07B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

WPI Data, PAJ, CHEM ABS Data, BIOSIS, MEDLINE, EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	WO 97 18471 A (ABBOTT LABORATORIES) 22 May 1997 (1997-05-22) claim 1 ---	1-25
Y	WO 97 14814 A (SMITHKLINE BEECHAM CORPORATION) 24 April 1997 (1997-04-24) claim 1 --- -/--	1-25

[X] Further documents are listed in the continuation of box C.   [X] Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

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"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
19 July 2000	31/07/2000

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Van Bohemen, C

Form PCT/ISA/210 (second sheet) (July 1992)

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication,where appropriate, of the relevant passages	Relevant to claim No.
X,P	CHEMICAL ABSTRACTS, vol. 132, Columbus, Ohio, US; abstract no. 175290, XP002142945 abstract & A. MEDEK ET AL.: "The Use of differential chemical shifts for determining the binding site location and orientation of protein-bound ligands" JOURNAL OF THE AMERICAN CHEMICAL SOCIETY , vol. 122, no. 6, 1 June 2000 (2000-06-01), pages 1241-1242, Washington DC USA	1-25

2

INTERNATIONAL SEARCH REPORT

		Inter)nal Application No
		PCT/US 00/07524

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 9718471	A	22-05-1997	US 5698401 A	16-12-1997
			AT 189063 T	15-02-2000
			AU 701810 B	04-02-1999
			AU 7732896 A	05-06-1997
			DE 69606324 D	24-02-2000
			DE 69606324 T	06-07-2000
			EP 0866967 A	30-09-1998
			EP 0981049 A	23-02-2000
			ES 2143793 T	16-05-2000
			JP 3032301 B	17-04-2000
			JP 11513498 T	16-11-1999
			PT 866967 T	28-04-2000
			US 5804390 A	08-09-1998
WO 9714814	A	24-04-1997	EP 0856067 A	05-08-1998
			JP 2000500565 T	18-01-2000

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(54) Title: USE OF 13C-NMR TO DETECT BINDING
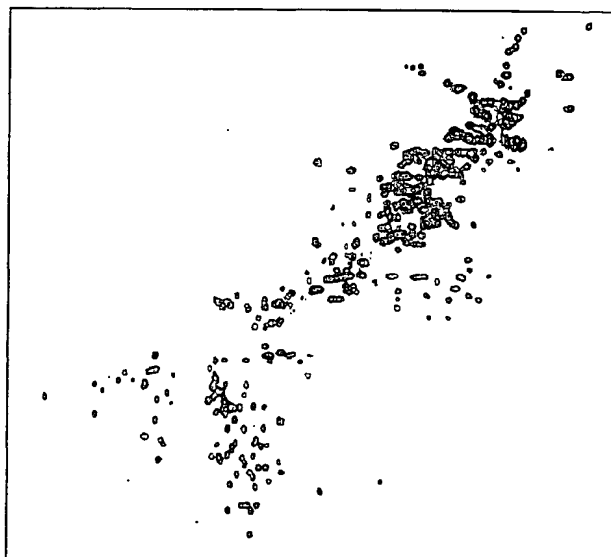
(57) Abstract: Methods of detecting binding of a putative ligand to a 13C-enriched target molecule, methods of screening for compounds which bind to a 13C-enriched target molecule, methods for calculating the dissociation constant of a ligand compound which binds to a 13C-enriched target molecule, and methods employed in the determination of the specific amino acids in a 13C-enriched target molecule affected by the binding of a ligand, as well as compounds identified by these screening methods, are provided herewith.

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